

PATENT COOPERATION TREATY

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NOTIFICATION OF ELECTION

(PCT Rule 61.2)

From the INTERNATIONAL BUREAU

To:

Commissioner
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 in its capacity as elected Office

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Applicant SELLERGREN, Börje et al	

1. The designated Office is hereby notified of its election made:



in the demand filed with the International Preliminary Examining Authority on:

22 March 2001 (22.03.01)



in a notice effecting later election filed with the International Bureau on:

2. The election ☒ was

was not

made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland Facsimile No.: (41-22) 740.14.35	Authorized officer Claudio Borton Telephone No.: (41-22) 338.83.38
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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

REC'D 24 OCT 2001

WIPO

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14

Applicant's or agent's file reference PC-2001870	FOR FURTHER ACTION See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)	
International application No. PCT/SE00/01776	International filing date (day/month/year) 14.09.2000	Priority date (day/month/year) 17.09.1999
International Patent Classification (IPC) or national classification and IPC ⁷ C08F 291/00, C08F 285/00, C08F 292/00		
Applicant MIP Technologies AB et al		

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.
2. This REPORT consists of a total of 4 sheets, including this cover sheet.
- ☒ This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

These annexes consist of a total of 4 sheets.

3. This report contains indications relating to the following items:

- I ☒ Basis of the report
- II ☐ Priority
- III ☐ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
- IV ☐ Lack of unity of invention
- V ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- VI ☐ Certain documents cited
- VII ☐ Certain defects in the international application
- VIII ☐ Certain observations on the international application

Date of submission of the demand 22.03.2001	Date of completion of this report 19.09.2001
Name and mailing address of the IPEA/SE Patent- och registreringsverket Box 5055 S-102 42 STOCKHOLM Facsimile No. 08-667 72 88	Authorized officer Monika Bohlin/EK Telephone No. 08-782 25 00

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.

PCT/SE00/01776

I. Basis of the report**1. With regard to the elements of the international application:***

- ☐ the international application as originally filed
- ☒ the description:
pages 1-15, as originally filed
pages _____, filed with the demand
pages _____, filed with the letter of _____
- ☒ the claims:
pages _____, as originally filed
pages _____, as amended (together with any statement) under article 19
pages 1-4, filed with the demand
pages _____, filed with the letter of _____
- ☒ the drawings:
pages 1-5, as originally filed
pages _____, filed with the demand
pages _____, filed with the letter of _____
- ☐ the sequence listing part of the description:
pages _____, as originally filed
pages _____, filed with the demand
pages _____, filed with the letter of _____

2. With regard to the language, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.
These elements were available or furnished to this Authority in the following language _____ which is:

- ☐ the language of a translation furnished for the purposes of international search (under Rule 23.1(b)).
- ☐ the language of publication of the international application (under Rule 48.3(b)).
- ☐ the language of the translation furnished for the purposes of international preliminary examination (under Rules 55.2 and/or 55.3).

3. With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- ☐ contained in the international application in written form.
- ☐ filed together with the international application in computer readable form.
- ☐ furnished subsequently to this Authority in written form.
- ☐ furnished subsequently to this Authority in computer readable form.
- ☐ The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
- ☐ The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. ☐ The amendments have resulted in the cancellation of:

- ☐ the description, pages _____
- ☐ the claims, Nos. _____
- ☐ the drawings, sheet/fig _____

5. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2 (c)).**

* Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are annexed to this report since they do not contain amendments (Rules 70.16 and 70.17).

** Any replacement sheet containing such amendments must be referred to under item I and annexed to this report.

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement**1. Statement**

Novelty (N)	Claims	<u>1-23</u>	YES
	Claims		NO
Inventive step (IS)	Claims	<u>1-23</u>	YES
	Claims		NO
Industrial applicability (IA)	Claims	<u>1-23</u>	YES
	Claims		NO

2. Citations and explanations (Rule 70.7)

This preliminary examination report is based on the claims 1-23 amended under Article 34 PCT and filed with the demand.

The following documents were cited in the search report:

- D1 Langmuir, Volume 15, 1999, Y. Nakayama et al
D2 J. Chem. Tech. Biotechnol., Volume 70, 1997, Hong Ying Wang et al

D1 relates to surface modification by sequentially forming polymer blocks on a polystyrene surface for biomedical applications, such as heparin immobilisation, protein immobilisation and drug release. According to D1 polymerisation on the substrate surface is conducted first, and then the bioactive substance (template) is attached. D1 does not disclose that a template molecule is first associated with a functional monomer, which is then polymerised in presence of a crosslinking monomer to provide a molecularly imprinted polymer (MIP).

D2 discloses molecular imprinting of N,N'-diethylaminodithiocarbamoylmethylstyrene (DTCS)-modified polyacrylonitrile membranes using theophylline as a template molecule in the polymerisation of acrylic acid and N,N'-methylenebisacrylamide on the surface of the membrane.

None of the cited documents discloses that the supported MIP is separated from the polymerisation medium, which is then reused for repeated preparation of MIP:s. This reuse is new, it improves the yield of the MIP and lowers the consumption of template and functional monomer.

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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.

PCT/SE00/01776

Supplemental Box

(To be used when the space in any of the preceding boxes is not sufficient)

Continuation of: Box V.

The invention according to the amended claims 1-23 is, in view of the arguments presented above, considered to be novel, to involve an inventive step and to be industrially applicable.

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(71) Applicant (for all designated States except US): MIP
TECHNOLOGIES AB [SE/SE]; Research Park Ideon,
S-223 70 Lund (SE).

(72) Inventors; and

(75) Inventors/Applicants (for US only): SELLERGREN,
Börje [SE/DE]; University of Mainz, Department of
Inorganic Chemistry and Analytical Chemistry, Duesberg-
weg 10-14, 55099 Mainz (DE). SULITZKY, Claudia
[DE/DE]; University of Mainz, Department of Inorganic

Chemistry and Analytical Chemistry, Duesbergweg 10-14,
55099 Mainz (DE). RÜCKERT, Bärbel [DE/DE]; Uni-
versity of Mainz, Department of Inorganic Chemistry and
Analytical Chemistry, Duesbergweg 10-14, 55099 Mainz
(DE).

(74) Agent: AWAPATENT AB; Box 5117, S-200 71 Malmö
(SE).

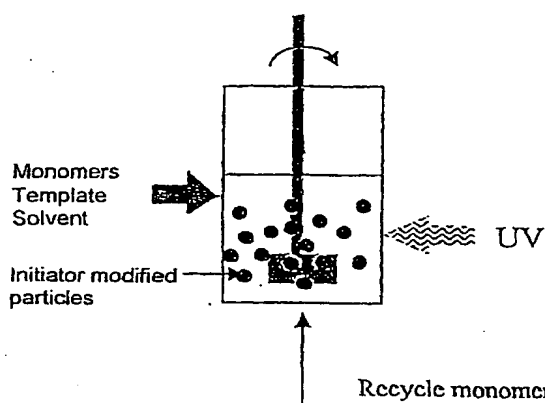
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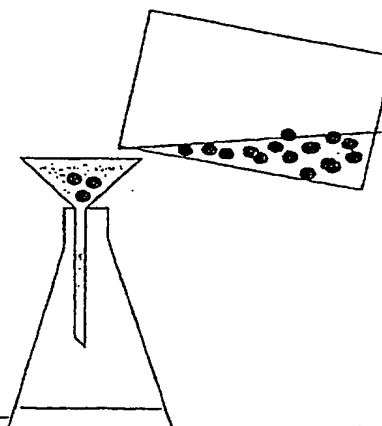
[Continued on next page]

(54) Title: NEW MOLECULARLY IMPRINTED POLYMERS GRAFTED ON SOLID SUPPORTS

A. Graft polymerization



B. Filter, wash, and dry
the particles



(57) Abstract: The invention refers to a molecularly imprinted polymer, a method of preparing a molecularly imprinted polymer material, and the use thereof. According to the invention a support and a composition comprising at least one monomer, and a template, in a polymerisation medium is polymerised with a free radical initiator, whereafter the template is removed from the molecularly imprinted polymer obtained. The polymerisation is confined to the surface of the support, preferably by confining the free radical initiator to the support by bonding or adsorption. The molecularly imprinted polymer may be used in chromatography, for separations, in chemical sensors, in molecular recognition as stationary phase in capillaries, in selective sample enrichment or in catalysis.



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- Before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments.

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

NEW MOLECULARLY IMPRINTED POLYMERS GRAFTED ON SOLID
SUPPORTS

Technical Field of the Invention

The present invention relates to a molecularly imprinted polymer, to a method for preparing said molecularly imprinted polymer, and to the use of said molecularly imprinted polymer.

Background Art

In the fields of medical, dietary, environmental and chemical sciences there is an increasing need for the selective separation of specific substances in complex mixtures of related substances. The end goal can be the preparative isolation of a certain compound or compounds or measurements of their concentration. Molecularly imprinted polymers (MIPs) often exhibit a high selectivity towards their substrate in analogy with the antibody-antigen complementarity. (1, 2) The technique shows promise in chiral separations of for example amino acid derivatives, peptides, phosphonates, aminoalcohols and beta-blocking compounds, affinity chromatography of nucleotides and the DNA-bases as well as substitute for antibodies in immunoassays for commercial drugs. Molecular imprinting (MI) consists of the following key steps: (1) Functional monomers are allowed to interact reversibly with a template molecule in solution. (2) The hereby formed template assemblies are copolymerised with a cross-linking monomer resulting in a cross-linked network polymer. (3) The template is displaced and the resulting MIP material can be used for selective molecular recognition of the corresponding compound. If the MIP material is crushed and sieved it can be packed in a chromatographic column and used for chromatographic separation of the template from structurally related analogs. Analytical as well as preparative applications are here possible. Preparative applications can be separation

of a compound from a complex mixture of structurally related compounds and isolation of the compound. This can be through an affinity chromatographic procedure where pH, ion strength or solvent gradients can be used in order to control the strength of interaction with the stationary phase. The separation can target enantiomers or diastereomers in a mixture of enantiomers or diastereomers of one or many compounds. Analytical applications can in addition to the above mentioned separations be: competitive binding assays, chemical sensors or selective sample enrichments.

Currently the most widely applied technique to generate molecularly imprinted binding sites is represented by the non-covalent route developed by the group of Mosbach(3). This makes use of non-covalent self-assembly of the template with functional monomers prior to polymerisation, free radical polymerisation with a cross-linking monomer and then template extraction followed by re-binding by non-covalent interactions. Although the preparation of a MIP by this method is technically simple it relies on the success of stabilisation of the relatively weak interactions between the template and the functional monomers. Stable monomer-template assemblies will in turn lead to a larger concentration of high affinity binding sites in the resulting polymer. The materials can be synthesized in any standard equipped laboratory in a relatively short time and some of the MIPs exhibit binding affinities and selectivities in the order of those exhibited by antibodies towards their antigens. Most MIPs are synthesized by free radical polymerisation of functional monounsaturated (vinyllic, acrylic, methacrylic) monomers and an excess of cross-linking di- or triunsaturated (vinyllic, acrylic, methacrylic) monomers resulting in porous organic network materials. These polymerisations have the advantage of being relatively robust allowing polymers to be prepared in high yield using different solvents (aqueous or organic) and at different

temperatures (4).. This is necessary in view of the varying solubilities of the template molecules.

The most successful non-covalent imprinting systems are based on commodity acrylic or methacrylic monomers, such as methacrylic acid (MAA), cross-linked with ethyleneglycol dimethacrylate (EDMA). Initially, derivatives of amino acid enantiomers were used as templates for the preparation of imprinted stationary phases for chiral separations (MICSPs) but this system has proven generally applicable to the imprinting of templates allowing hydrogen bonding or electrostatic interactions to develop with MAA. (5, 6) The procedure applied to the imprinting with L-phenylalanine anilide (L-PA) is outlined in Fig. 1. In the first step, the template (L-PA), the functional monomer (MAA) and the cross-linking monomer (EDMA) are dissolved in a poorly hydrogen bonding solvent (diluent) of low to medium polarity. The free radical polymerisation is then initiated with an azo initiator, commonly azo-N,N'-bis-isobutyronitrile (AIBN) either by photochemical homolysis below room temperature (6, 7) or thermochemically at 60°C or higher (5). Lower thermochemical initiation temperatures down to 40°C or 30°C may be obtained using azo-N,N'-bis-divaleronitrile (ABDV) and V70 resp. instead of AIBN as initiator (see). (7, 8) In the final step, the resultant polymer is crushed by mortar and pestle or in a ball mill, extracted by a Soxhlet apparatus, and sieved to a particle size suitable for chromatographic (25-38 μm) or batch (150-250 μm) applications. (6) The polymers are then evaluated as stationary phases in chromatography by comparing the retention time or capacity factor (k') (9) of the template with that of structurally related analogs.

As appears from above MIPs have so far been prepared in the form of continuous blocks that need to be crushed and sieved before use. This results in a low yield of irregular particles, a high consumption of template and a material exhibiting low chromatographic efficiency. There

is therefore a need for MI-materials that can be prepared in high yield in the form of regularly shaped particles with low size dispersity and a controlled porosity. These are expected to be superior in terms of mass transfer characteristics and sample load capacity compared to the materials obtained from the monolithic approach.

Such MIPs have been previously prepared through suspension(10, 11)- polymerisation techniques, dispersion polymerisation(12) or precipitation polymerisation(13). This resulted in spherical particles of a narrow size distribution. These procedures have the limitation of being very sensitive to small changes in the manufacturing conditions and the type of solvents and polymerisation conditions that can be applied. Thus the procedures need careful optimization for each new template target which significantly reduces the usefulness of this route. Moreover conditions leading to low dispersity spherical particles may not be compatible with conditions leading to high selectivity and affinity for the template target.

An alternative to this procedure is the coating of preformed support materials.(14-16) MIPs have been prepared as grafted coatings on oxide supports(14, 16) on organic polymer supports(15) and on the walls of fused silica capillaries(17-19). The former technique allows the use of the wide variety of oxide support materials available with different sizes and porosities. Grafting techniques to prepare organic polymer coatings are expected to be generally applicable to molecular imprinting since the structure of the underlying support is already fixed. Thus compared to the large number of factors influencing the end result in suspension or precipitation type polymerisations a smaller number of factors is likely to influence the end result in the preparation of the imprinted coatings. This will make the grafted coatings techniques less sensitive to changes in conditions offering a more robust method. These types of coating techniques are furthermore applicable to modify surfaces of

monolithic type supports or microchips prepared by lithographic techniques. The oxide based materials are rigid porous supports with a limited inner pore volume. An alternative support that could potentially carry more grafted imprinted polymer per unit weight and thus allow a higher density of imprinted sites would be to make use of swellable organic resins. In this context Merrifield resins containing grafted initiator or monomer could be used.

10 Sofar most imprinted coatings have been prepared by grafting polymers to the various surfaces. Thus the surface contains prior to polymerisation polymerizable double bonds that can add to the growing polymer chains in solution linking them to the surface. The problem with
15 this technique is the presence of initiator in solution requiring the monomer mixture to be applied as a liquid thin film on the surface prior to polymerisation. Thus the exact amount of monomers that will coat the available surface with an up to ca 100 Å thick liquid film is dissolved together with initiator in an excess of solvent.
20 Thereafter the modified support is added and the solvent evaporated to leave the monomer film and initiator on the surface. Polymerisation is then carried out usually at elevated temperatures. With this procedure the thickness
25 of the polymer layer is difficult to control and capillary forces upon evaporation of solvent may cause incomplete wetting of the surface. Moreover a continuous method of synthesising the particles is difficult to envisage with this method.

30 A considerable improvement in this regard would be to confine the initiator radicals to the support surface (Fig. 2). (20, 21) In absence of chain transfer this would lead to chain growth occurring only from the surface of the support with no polymerisation occurring in solution.
35 For molecular imprinting this would have important consequences. For instance the polymerisation can be carried out on the surface of initiator modified support particl-

es suspended in a mixture of the monomers and solvent. This would allow polymerisation in a simple tank reactor by either thermal or photochemical initiation. The latter technique would allow the particles to be modified during
5 the sedimentation possibly leading to a continuous method for preparing the imprinted composite particles (Fig. 3). Polymerisation would here only occur on the particle surface leaving the solution containing the monomers unreacted. The monomer solution can thus be reused for the
10 coating of several batches of particles. The problem of confining polymer chain growth to the support surface and suppress it in solution can be solved by attaching the radical initiator so that the radical formed upon bond homolysis remains bound to the surface. Alternatively the
15 radical formed that is not attached to the surface should undergo rapid reaction to give an unreactive species. It should be possible to prepare the grafted coatings using monomers such as those based on styren/divinylbenzene, methacrylates, acrylates, acrylamides and in the presence
20 of one or more template molecules.

Summary of the Invention

Thus, the present invention relates to a molecularly imprinted polymer obtainable by polymerising a composition comprising at least one monomer, and a template, on
25 a support in a polymerisation medium with a free radical initiator, whereafter the template is removed from the molecularly imprinted polymer obtained, said polymerisation being confined to the surface of the support.

The invention further relates to a method for preparing a molecularly imprinted polymer which comprises polymerising a composition comprising at least one monomer, and a template, on a support in a polymerisation medium with a free radical initiator, whereafter the template is removed from the molecularly imprinted polymer obtained, said polymerisation being confined to the surface of the support.

Still further the invention relates to the use of a molecularly imprinted polymer as defined above in chromatography, for separations, in chemical sensors, in molecular recognition as stationary phase in capillaries, in selective sample enrichment or in catalysis.

These and other advantages and characterising features of the present invention will appear from the following specification and the appended claims.

Brief Description of the Drawings

Fig. 1 illustrates molecular imprinting with L-phenylalanine anilide (L-PA).

Fig. 2 illustrates the procedure of confining initiator radicals to the surface of a support.

Fig. 3 illustrates a method for preparing imprinted composite particles.

Fig. 4A illustrates the use of a presynthesized azosilane initiator where both ends may be attached to the surface of a support.

Fig. 4B illustrates an initiator that may be preadsorbed on a support surface and that is insoluble in the monomer containing solution.

Fig. 4C illustrates the use of microwaves to selectively heat the particle surface.

Fig. 4D illustrates the use of iniferters such as dithiocarbamate coupled onto the surface.

Detailed Description of the Invention

The invention will now be described in more detail with reference to a number of non-limiting examples:

The invention refers to a material that consists in a support (porous or nonporous material or planar surface) coated with a polymer layer, a method for its fabrication and use of said material in for instance chromatography, for separations, in chemical sensors, in selective sample enrichment, in molecular recognition as stationary phase in capillaries or in catalysis. The material is prepared by grafting a polymer layer on the surface of a preformed organic or inorganic support material or surface. The grafting can be combined with the technique of molecular imprinting.

In one embodiment of the present invention the polymerisation is confined to the surface of the support by confining the free radical initiator to the support. According to one aspect the free radical initiator is bound (covalently or non-covalently such as e.g. by hydrogen bonds) to the surface of the support. According to another aspect the free radical initiator is adsorbed to the surface of the support, preferably by dissolving it in a solvent for the free radical initiator, applying the solution to the support, and removing the solvent, said free radical initiator being insoluble in the polymerisation medium or remaining attached to the support surface by adsorptive forces.

In another embodiment of the present invention the polymerisation is confined to the surface of the support by subjecting the composition, the support and the free radical initiator to microwave irradiation which selectively heats the support and thereby initiates a polymerisation reaction at the surface of the support.

In a further embodiment of the present invention the polymerisation is repeated at least once with a different composition to obtain at least one further layer of molecularly imprinted polymer. This allows the manufacturing of layered surfaces containing one or more imprinted layers using possibly different templates and layers of different polarity or other functional properties.

The support used in the present invention is preferably selected from the group consisting of porous and non-porous, planar and non-planar inorganic and organic supports. As examples of such support materials may be mentioned oxides such as alumina and silica, and organic resins in the form of particles such as spheres, or sheets.

The template used in the present invention may be any molecule or ion and is preferably selected from the group consisting of organic or inorganic molecule entities, ions, antibodies, antigens, amino acids, peptides, proteins, nucleotides, DNA-bases, carbohydrates, drugs, pesticides, and derivatives thereof, etc.

The expression "polymerisation medium" as used herein means a liquid medium in which the polymerisation is carried out. The polymerisation medium may e.g. be a solvent in which the monomers are soluble. It may also be a monomer acting as a solvent for the other components of the polymerisable composition.

The support surface is prepared as follows. A free radical initiator is bound to the surface either covalently or noncovalently so that the free radicals generated upon initiation remain confined to the surface or vicinity of the surface. The absence of polymer propagation in solution will lead to a higher accessibility of the monomers at the surface. Furthermore this method will allow the tuning of the thickness of the polymer layer.

Surface attachment of a free radical initiator has been disclosed generally by Guyot et.al. (21) and Tsubokawa et.al. (22, 23) It relies on presilanization of the surface using 3-aminopropyltriethoxysilane or a glycidoxypropylsilane (GPS) followed by reaction of the amino groups or the epoxy groups with an azoinitiator such as azo-bis(cyanopentanoic acid, ACPA) leading to the formation of an amide (using DCC as condensing reagent) or ester link between the surface and the azoinitiator. Also peroxy initiators may be used although better re-

sults are obtained using the grafted azoinitiator followed by photochemical initiation. High yields of grafted polymer are obtained using silica reacted with toluene-2,4-diisocyanate (TDI) followed by reaction with ACPA.

5 Example 1

Coupling of initiator to amino, epoxy or chloromethyl modified supports or resins

Epoxy and chloromethyl modified supports: A typical example is as follows. Into a flask, 3 g of epoxy modified particles 50 mL of DMSO, 0.5 g of ACPA and picoline were charged. The reaction mixture was stirred for 5 h at 50°C. After the reaction the particles were washed with methanol and dried.

Amino modified supports: A typical example is as follows. Into a flask, 3 g of epoxy modified particles 50 mL of DMF, 0.5 g of ACPA and dicyclohexyldicarbodiimide (DCCI) and base were charged. The reaction mixture was stirred for 5 h. After the reaction the particles were washed with methanol and dried.

20 The above procedure does not confine all initiator radicals to the surface since the initiator is bound at only one position. This invention describes three alternative procedures to confine the polymerisation to the surface.

25 1. The use of a presynthesized azosilane (Fig. 4A). This will more likely lead to a two point attachment of the initiator to the surface.

Example 2

30 Synthesis of azosilane for two point coupling of an azoinitiator to a surface or support

The azosilane was synthesized by mixing 0.5 mole glycidoxypropyltrimethoxysilane (GPS) and 0.25 mole ACPA in 200 mL isopropanol and catalytic amounts of picoline. The reaction was allowed to continue at room temperature and the product isolated by evaporation to dryness followed by purification by column chromatography giving the product in 60 % yield.

Example 3

Coupling of silane to a surface

5 The silane was coupled by reaction in water at low temperature (20°C) for 24 hours.

2. Preadsorbtion of an initiator that is insoluble in the monomer containing solution. Thus, a polar water soluble initiator as for instance an azo-bis-amidine, (24) can be adsorbed to the surface from aqueous solvent, the surface dried and then the polymerisation initiated as described above (Fig. 4B). The free radicals generated from the initiator will stay associated to the surface due to their insolubility in the monomer mixture.

15 Example 4

Adsorption of amidineazoinitiator to a support surface

 An amidineazoinitiator such as 2,2'-azobis(N,N'-dimethyleneisobutyramidine) or 2,2'-azobis(2-amidino-propane) is dissolved in methanol/water and support particles such as silica are added. After several hours of equilibration the solvent is removed by filtration and the particles dried under vacuum.

3. Use of microwaves to selectively heat the particle surface (Fig. 4C).

25 Example 5

Microwave initiated polymerisation

 Particles are added to a solution of monomers and initiator in a suitable solvent. The polymerisation is initiated by microwave irradiation at a wavelength causing local heating of the particles only.

4. Use of iniferters such as dithiocarbamate coupled onto the surface (Fig. 4D). (25) (The term "iniferter" is an abbreviation for "initiator + transfer agent + terminator").

Example 6

Synthesis of support or polymer-bound initiator

To a surface or polymer containing bound chloromethyl groups is given N,N-diethyldithiocarbamate in solution and the reaction allowed to proceed at elevated temperatures.

Example 7

Synthesis of block-graft imprinted copolymer

Particles or a surface containing bound dithiocarbamate groups are/is added to a mixture of monomers (concentration about 5 moles/litre), template and solvent under nitrogen. The polymerisation was initiated by irradiation with an ultrahigh pressure mercury UV lamp and allowed to proceed for a certain time. Then the unreacted monomers and template were washed away. The obtained particles or surface can then be immersed in another solution containing another monomer and the procedure repeated. This allows the manufacturing of layered surfaces containing one or more imprinted layers using possibly different templates and layers of different polarity or other functional properties.

Example 8

Endcapping of unreacted silanol groups

Prior to polymerisation endcapping of unreacted silanol groups can be done. Hexamethylsilazane is here effective. Good wetting is critical for the formation of a homogenous layer fully covering the support. Another possibility to enhance the wetting is to use organosilanes containing functionalities resembling solvents known to be good solvents for the methacrylate polymerisations. Among these chlorinated hydrocarbons are particularly useful.

Grafting of polymer layer

The polymerisation can be carried out in a stirred suspension of the particles in the monomer mixture since growth only takes place on the surface (see Fig. 3). Thus the initiator modified particles are added to a

monomer containing solution and solvent and possibly a template and the suspension stirred. The polymerisation is then carried out photochemically or thermally. The particles can be based on any inorganic or organic support material and the template on any molecule or ion dissolved in the monomer mixture solution. The grafting can also occur on other surfaces such as those generated by lithographic processes or on the walls of capillaries or fibres. The thickness of the polymer layer is tunable by varying the time of reaction.

Example

To a stirred solution of 38 ml (0.2 mole) EDMA, 3.4 ml (40 mmole) MAA and 10 mmole terbutylazine (or no template) in 56 ml dichloromethane is added 5 g of any of the initiator modified particles described in Examples 1-6. The suspension is sparged with nitrogen and the polymerisation initiated by UV irradiation using a standard high pressure mercury lamp at 15°C or by heating to a temperature providing a suitable rate of polymerisation. The suspension is stirred under nitrogen and UV irradiation or heating for 24 h and the particles then filtered, washed and dried under vacuum. The monomer mixture is then used to modify a second batch of particles.

The resulting particles exhibit high selectivity and affinity for the template, terbutylazine.

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CLAIMS

1. A molecularly imprinted polymer, c h a r a c -
t e r i s e d in that it is obtainable by polymerising a
5 composition comprising at least one monomer, and a tem-
plate, on a support in a polymerisation medium with a
free radical initiator, whereafter the template is re-
moved from the molecularly imprinted polymer obtained,
and that said polymerisation is confined to the surface
10 of the support.

2. A molecularly imprinted polymer according to
claim 1, wherein the polymerisation is confined to the
surface of the support by confining the free radical ini-
tiator to the support.

15 3. A molecularly imprinted polymer according to
claim 2, wherein the free radical initiator is bound or
adsorbed to the surface of the support.

4. A molecularly imprinted polymer according to
claim 1, wherein the polymerisation is confined to the
20 surface of the support by subjecting the composition, the
support and the free radical initiator to microwave irra-
diation which selectively heats the support and thereby
initiates a polymerisation reaction at the surface of the
support.

25 5. A molecularly imprinted polymer according to any
one of claims 1-4, wherein the polymerisation is repeated
at least once with a different composition to obtain at
least one further layer of molecularly imprinted polymer.

30 6. A molecularly imprinted polymer according to any
one of claims 1-5, wherein the support is selected from
the group consisting of porous and non-porous, planar and
non-planar inorganic and organic supports.

7. A molecularly imprinted polymer according to any
one of claims 1-6, wherein the template is selected from
35 the group consisting of organic or inorganic molecule en-
tities, ions, antibodies, antigens, amino acids, pep-

tides, proteins, nucleotides, DNA-bases, carbohydrates, drugs, pesticides, and derivatives thereof.

8. A method for preparing a molecularly imprinted polymer c h a r a c t e r i s e d by polymerising a composition comprising at least one monomer, and a template, on a support in a polymerisation medium with a free radical initiator, whereafter the template is removed from the molecularly imprinted polymer obtained, said polymerisation being confined to the surface of the support.

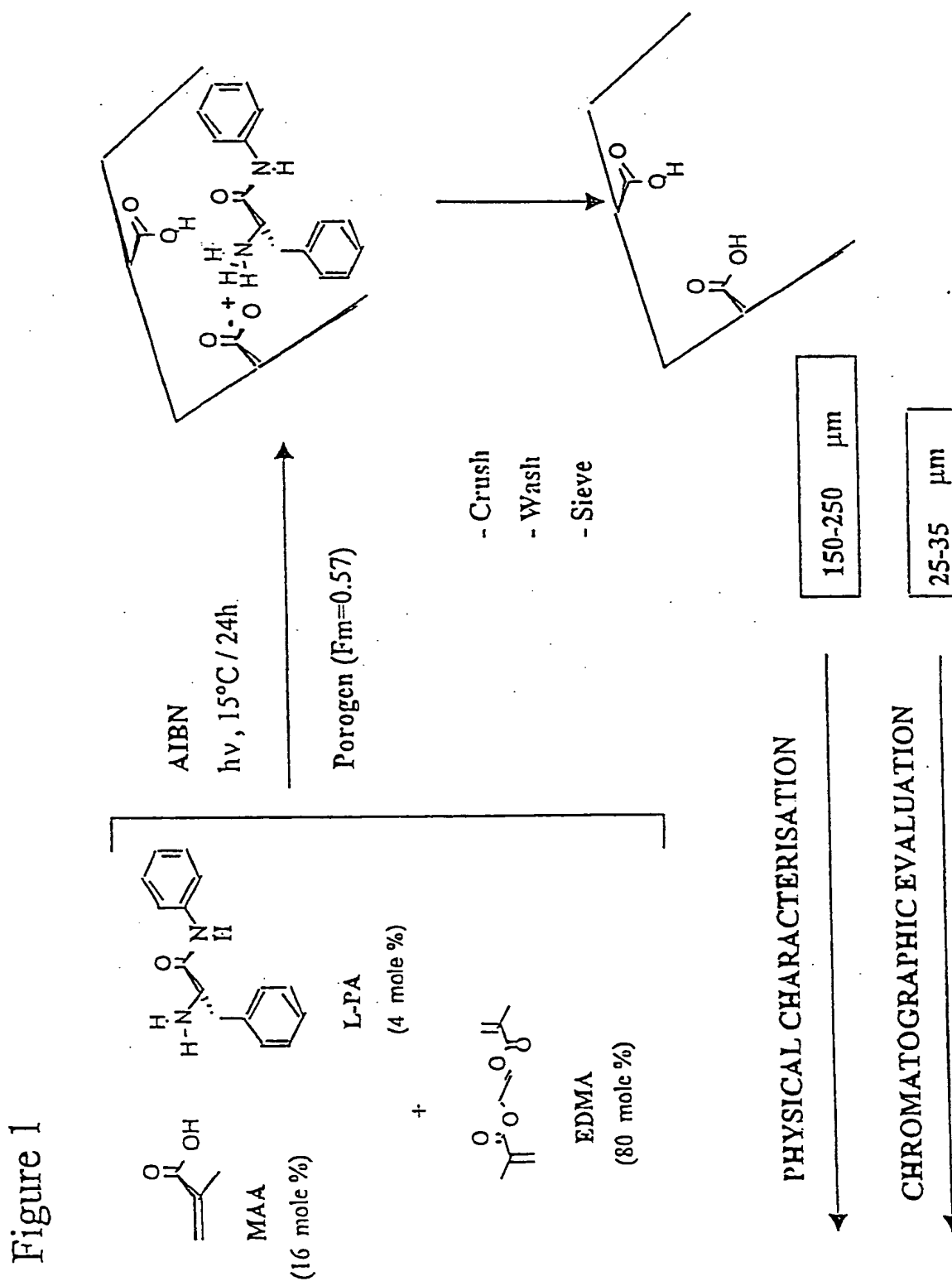
9. A method according to claim 8, wherein the polymerisation is confined to the surface of the support by confining the free radical initiator to the support.

10. A method according to claim 9, wherein the free radical initiator is bound or adsorbed to the surface of the support.

11. A method according to claim 8, wherein the polymerisation is confined to the surface of the support by subjecting the composition, the support and the free radical initiator to microwave irradiation which selectively heats the support and thereby initiates a polymerisation reaction at the surface of the support.

12. A method according to any one of claims 8-11, wherein the polymerisation is repeated at least once with a different composition to obtain at least one further layer of molecularly imprinted polymer.

13. Use of a molecularly imprinted polymer according to any one of claims 1-7, or prepared according to any one of claims 8-13, in chromatography, for separations, in chemical sensors, in molecular recognition as stationary phase in capillaries, in selective sample enrichment or in catalysis.



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Figure 3

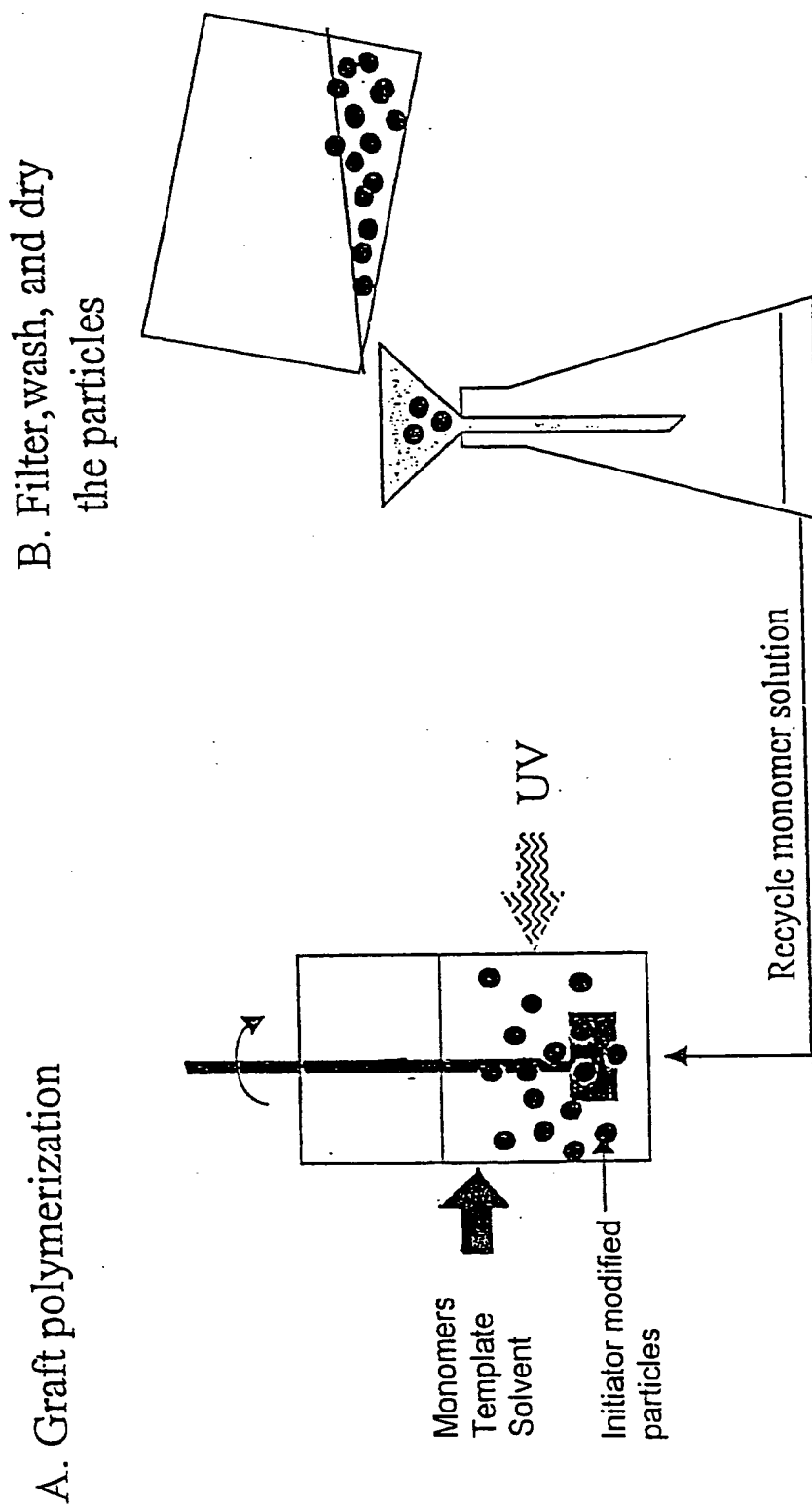
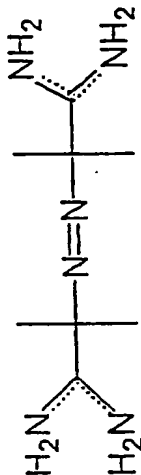
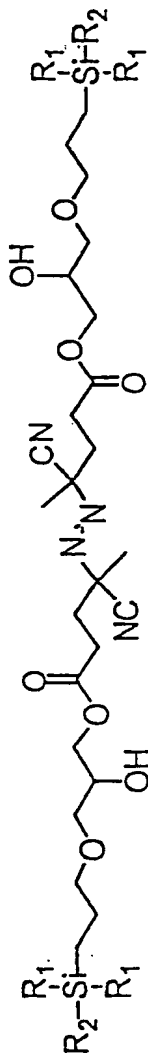
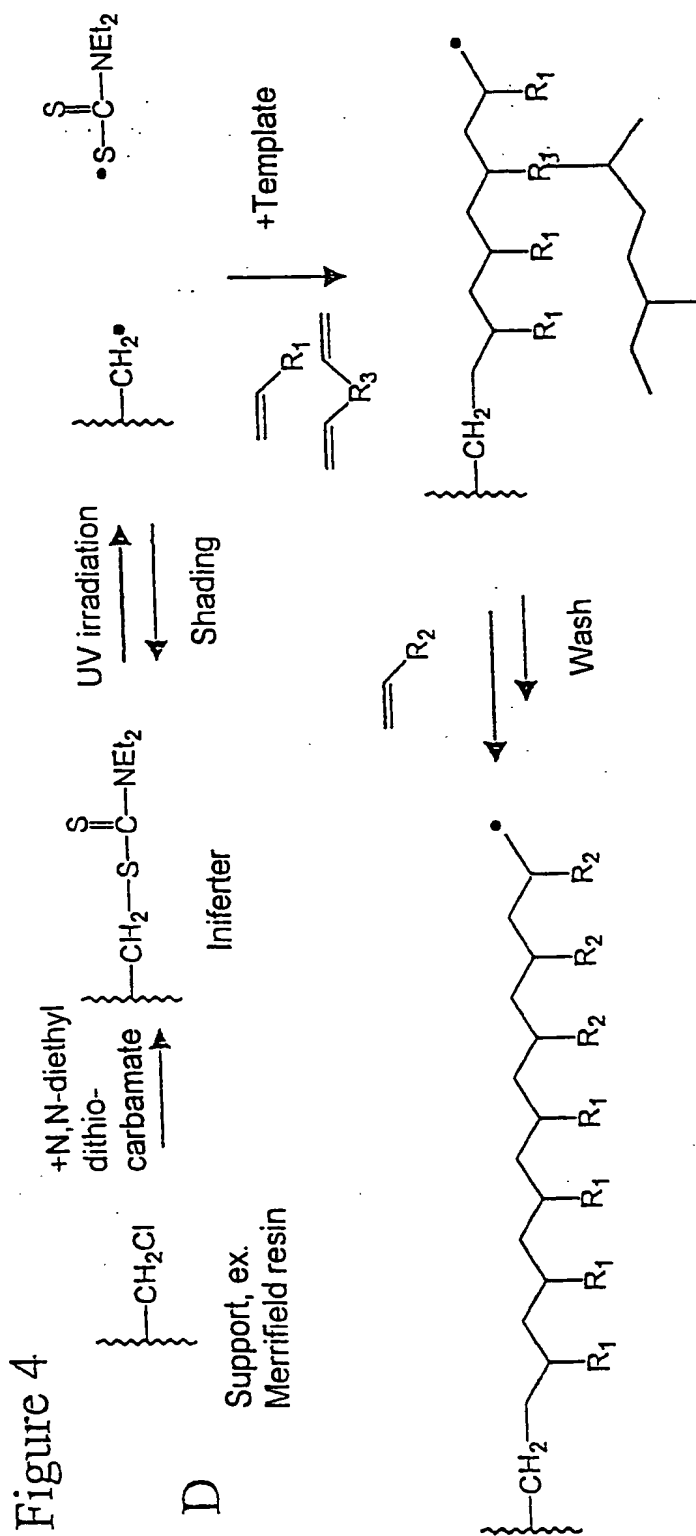


Figure 4



Microwave

C



INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 00/01776

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: C08F 291/00, C08F 285/00, C08F 292/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: C08F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Langmuir, Volume 15, 1999, Y. Nakayama et al, "Surface Macromolecular Microarchitecture Design: Biocompatible Surfaces via Photo-Block-Graft-Copolymerization Using N, N-Diethyldithiocarbamate" page 5560 - page 5566 --	1-3,5-10, 12-13
X	J. Chem. Tech. Biotechnol, Volume 70, 1997, Hong Ying Wang et al, "Surface Molecular Imprinting on Photosensitive Dithiocarbamoyl Polyacrylonitrile Membranes Using Photograft Polymerization" page 355 - page 362 --	1-3,5-10, 12-13

☒ Further documents are listed in the continuation of Box C.☒ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

12 January 2001

Date of mailing of the international search report

15-01-2001

Name and mailing address of the ISA/

Swedish Patent Office

Box 5055, S-102 42 STOCKHOLM

Facsimile No. +46 8 666 02 86

Authorized officer

Monika Bohlin/Els

Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 00/01776

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	Macromolecules, Volume 33, 2000, Sergey A. Piletsky et al, "Surface Functionalization of Porous Polypropylene Membranes with Molecularly Imprinted Polymers by Photograft Copolymerization in Water" page 3092 - page 3098 --	1-3,5-10, 12-13
P,A	WO 0007702 A2 (POLY-AN GMBH), 17 February 2000 (17.02.00), abstract --	1-13
A	Macromolecules, Volume 31, 1998, Liang Liang et al, "Reversible Surface Properties of Glass Plate and Capillary Tube Grafted by Photopolymerization of N-Isopropylacrylamide", page 7845 - page 7850, abstract -- -----	1-13

INTERNATIONAL SEARCH REPORT

Information on patent family members

04/12/00

International application No.

PCT/SE 00/01776

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 0007702 A2	17/02/00	DE 19936992 A	25/05/00

CLAIMS

1. A molecularly imprinted polymer, c h a r a c -
t e r i s e d in that it is obtainable by polymerising a
5 composition comprising at least one monomer, and a tem-
plate, on a support in a polymerisation medium with a
free radical initiator, whereafter the template is re-
moved from the molecularly imprinted polymer obtained,
and that said polymerisation is confined to the surface
10 of the support.

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surface of the support by confining the free radical ini-
tiator to the support.

15 3. A molecularly imprinted polymer according to
claim 2, wherein the free radical initiator is bound or
adsorbed to the surface of the support.

4. A molecularly imprinted polymer according to
claim 1, wherein the polymerisation is confined to the
20 surface of the support by subjecting the composition, the
support and the free radical initiator to microwave irra-
diation which selectively heats the support and thereby
initiates a polymerisation reaction at the surface of the
support.

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one of claims 1-4, wherein the polymerisation is repeated
at least once with a different composition to obtain at
least one further layer of molecularly imprinted polymer.

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one of claims 1-5, wherein the support is selected from
the group consisting of porous and non-porous, planar and
non-planar inorganic and organic supports.

35 7. A molecularly imprinted polymer according to any
one of claims 1-6, wherein the template is selected from
the group consisting of organic or inorganic molecule en-
tities, ions, antibodies, antigens, amino acids, pep-

tides, proteins, nucleotides, DNA-bases, carbohydrates, drugs, pesticides, and derivatives thereof.

8. A method for preparing a molecularly imprinted polymer c h a r a c t e r i s e d by polymerising a composition comprising at least one monomer, and a template, on a support in a polymerisation medium with a free radical initiator, whereafter the template is removed from the molecularly imprinted polymer obtained, said polymerisation being confined to the surface of the support.

9. A method according to claim 8, wherein the polymerisation is confined to the surface of the support by confining the free radical initiator to the support.

10. A method according to claim 9, wherein the free radical initiator is bound or adsorbed to the surface of the support.

11. A method according to claim 8, wherein the polymerisation is confined to the surface of the support by subjecting the composition, the support and the free radical initiator to microwave irradiation which selectively heats the support and thereby initiates a polymerisation reaction at the surface of the support.

12. A method according to any one of claims 8-11, wherein the polymerisation is repeated at least once with a different composition to obtain at least one further layer of molecularly imprinted polymer.

13. Use of a molecularly imprinted polymer according to any one of claims 1-7, or prepared according to any one of claims 8-13, in chromatography, for separations, in chemical sensors, in molecular recognition as stationary phase in capillaries, in selective sample enrichment or in catalysis.